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Novel peptidase inhibitors.

This invention relates to analogs of peptidase substrates in which the nitrogen atom of the scissile amide group of the substrate peptide has been replaced by a substituted malonyl moiety.

The contemplated peptidase inhibitors of the foregoing enzymes are selected from the generic formula

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 R_1NH C-X R_2 C

the hydrates, isosteres or the pharmaceutically acceptable salts thereof wherein X is

 R_1 is hydrogen, an amino protecting group selected from Group K, an α -amino acid or a peptide comprised of a number of α -amino acid building blocks, said α -amino acid or peptide optionally bearing on its terminal nitrogen atom an amino protecting group selected from Group K,

 R_2 is the "R group" residue of the α -amino acid responsible for directing the inhibitor to the active site of the enzyme or is -A-SiR₇R₈R₉, C₁₋₁₀ alkyl, aralkyl or aryl with R₇, R₈ and R₉, each being selected from C₁₋₁₀ alkyl, aralkyl or aryl and A is a C₁₋₆ alkylene.

 R_4 is the specific R-group residue of the $\alpha\text{-amino}$ acid for that peptidase substrate analog,

R₅ is an α-amino acid or peptide comprised of α-



EUROPEAN SEARCH REPORT



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tegory		with indication, where appropriate, elevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CLS)
X	EP-A-0 133 225 (HOFFI Example 5; pages 28-29		3	C 07 K 5/02. C 07 K 5/06 C 07 K 5/08
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				TECHNICAL FIELDS SEARCHED (Int. CI.5)
				C 07 K C 07 C A 61 K
	The present search report has	1990 drawn up for all claims	-	
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Y: par doo A: tec	CATEGORY OF CITED DOCUMENTS X: particularly relevant if taken alone Y: particularly relevant if combined with another document of the same catagory A: technological background O: non-written disclosure P: intermediate document		E: earlier patent document, but published on, or after the filing date D: document cited in the application L: document cited for other reasons L: member of the same patent family, corresponding	